

=> d his

(FILE 'HOME' ENTERED AT 15:55:08 ON 10 JUN 2003)

FILE 'MEDLINE' ENTERED AT 15:55:37 ON 10 JUN 2003

L1 13063 S SICKLE CELL
L2 39195 S ANTIVIRAL OR ACYCLOVIR OR VALACYCLOVIR
L3 318 S VALACYCLOVIR
L4 0 S L3 AND L1
L5 9 S L2 AND L1

FILE 'REGISTRY' ENTERED AT 16:00:41 ON 10 JUN 2003

L6 27 S ACYCLOVIR
L7 1 S ACYCLOVIR/CN
L8 1 S VALACYCLOVIR/CN

FILE 'MEDLINE' ENTERED AT 16:02:35 ON 10 JUN 2003

L9 5053 S 59277-89-3
L10 13070 S SICKLE# CELL
L11 2 S L10 AND L9
L12 0 S 124832-26-4

FILE 'REGISTRY' ENTERED AT 16:05:31 ON 10 JUN 2003

L13 1 S 124832-27-5

FILE 'BIOSIS' ENTERED AT 16:06:15 ON 10 JUN 2003

L14 406 S VALACYCLOVIR OR VALACICLOVIR
L15 5711 S ACYCLOVIR OR ZOVIRAX OR ACICLOVIR
L16 5853 S L15 OR L14
L17 10631 S SICKLE# CELL
L18 1 S L17 AND L16

FILE 'WPIDS' ENTERED AT 16:07:50 ON 10 JUN 2003

L19 11309 S ANTIVIRAL OR ACYCLOVIR OR VALACYCLOVIR
L20 30 S L19 AND L17

FILE 'USPATFULL' ENTERED AT 16:11:02 ON 10 JUN 2003

L21 3755 S ACYCLOVIR OR ACICLOVIR
L22 280 S VALACYCLOVIR OR VALACICLOVIR
L23 536 S 59277-89-3/RN
L24 66 S 124832-26-4/RN
L25 14 S 124832-27-5/RN
L26 3899 S L21 OR L22 OR L23 OR L24 OR L25
L27 3067 S SICKLE# CELL
L28 198 S L26 AND L27
L29 1 S L26 (P) L27

FILE 'MEDLINE' ENTERED AT 16:37:21 ON 10 JUN 2003

L30 6862 S HYDROXYUREA
L31 13070 S SICKLE# CELL
L32 324 S L30 AND L31
L33 12 S PREVENT SICKLING
L34 36259 S ANTIVIRAL
L35 2 S L34 AND L32
L36 0 S PD=1990
L37 0 S 1990/PD
L38 0 S 1990/DATE
L39 387715 S 1990/PY
L40 174 S L39 AND L30
L41 6 S L31 AND L40

=> log hold

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

	ENTRY	SESSION
FULL ESTIMATED COST	18.31	131.57

SESSION WILL BE HELD FOR 60 MINUTES
STN INTERNATIONAL SESSION SUSPENDED AT 16:58:57 ON 10 JUN 2003

=> d his

(FILE 'HOME' ENTERED AT 08:46:40 ON 11 JUN 2003)

FILE 'REGISTRY' ENTERED AT 08:47:10 ON 11 JUN 2003

L1 1 S 5-AZACYTIDINE/CN

FILE 'CA' ENTERED AT 08:47:57 ON 11 JUN 2003

L2 5088 S HYDROXYUREA

L3 37559 S ANTIVIRAL

L4 169 S L2 AND L3

L5 1401 S HYDROXYUREA/TI

L6 54 S L5 AND L3

S 320-67-2/REG#

FILE 'REGISTRY' ENTERED AT 08:54:27 ON 11 JUN 2003

L7 1 S 320-67-2/RN

FILE 'CA' ENTERED AT 08:54:27 ON 11 JUN 2003

L8 1266 S L7

L9 3408 S SICKLE# CELL

L10 14 S L8 AND L9

=> log hold

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

22.58

69.47

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE

TOTAL

ENTRY

SESSION

CA SUBSCRIBER PRICE

-2.48

-4.96

SESSION WILL BE HELD FOR 60 MINUTES

STN INTERNATIONAL SESSION SUSPENDED AT 09:01:10 ON 11 JUN 2003

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L6 ANSWER 51 OF 54 CA COPYRIGHT 2003 ACS
AN 121:292089 CA
TI **Hydroxyurea** as an inhibitor of human immunodeficiency virus-type
1 replication
AU Lori, Franco; Malykh, Andrei; Cara, Andrea; Sun, Daisy; Weinstein, John
N.; Lisziewicz, Julianna; Gallo, Robert C.
CS Laboratory of Tumor Cell Biology, National Institutes of Health, Bethesda,
MD, USA
SO Science (Washington, D. C.) (1994), 266(5186), 801-5
CODEN: SCIEAS; ISSN: 0036-8075
PB American Association for the Advancement of Science
DT Journal
LA English
AB Hydroxyurea, a drug widely used in therapy of several human diseases,
inhibits deoxynucleotide synthesis- and, consequently, DNA synthesis-by
blocking the cellular enzyme ribonucleotide reductase. Hydroxyurea
inhibits human immunodeficiency virus-type 1 (HIV-1) DNA synthesis in
activated peripheral blood lymphocytes by decreasing the amt. of
intracellular deoxynucleotides, thus suggesting that this drug has an
antiviral effect. Hydroxyurea has now been shown to block HIV-1
replication in acutely infected primary human lymphocytes (quiescent and
activated) and macrophages, as well as in blood cells infected in vivo
obtained from individuals with acquired immunodeficiency syndrome (AIDS).
The **antiviral** effect was achieved at nontoxic doses of
hydroxyurea, lower than those currently used in human therapy.
Combination of hydroxyurea with the nucleoside analog didanosine
(2',3'-dideoxyinosine, or ddI) generated a synergistic inhibitory effect
without increasing toxicity. In some instances, inhibition of HIV-1 by
hydroxyurea was irreversible, even several weeks after suspension of drug
treatment. The indirect inhibition of HIV-1 by hydroxyurea is not
expected to generate high rates of escape mutants. Hydroxyurea therefore
appears to be a possible candidate for AIDS therapy.

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